

MODIFIED BOVINE ADENOVIRUS HAVING ALTERED TROPISM

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This application claims the benefit of U.S. Provisional Application Serial No. 60/208,678, filed May 31, 2000, hereby incorporated herein in its entirety.

TECHNICAL FIELD

10 This invention relates to bovine adenoviruses comprising a modification in a capsid protein and which exhibit altered tropism. The present invention also relates to methods of making and using bovine adenoviruses having altered tropism.

BACKGROUND ART

15 The adenoviruses cause enteric or respiratory infection in humans as well as in domestic and laboratory animals. The bovine adenoviruses (BAV) comprise at least nine serotypes divided into two subgroups. These subgroups have been characterized based on enzyme-linked immunoassays (ELISA), serologic studies with immunofluorescence assays, virus-neutralization tests, immunoelectron microscopy, by their host specificity and clinical syndromes. Subgroup 1 viruses include BAV 1, 20 2, 3 and 9 and grow relatively well in established bovine cells compared to subgroup 2 which includes BAV 4, 5, 6, 7 and 8.

25 BAV3 was first isolated in 1965 and is the best characterized of the BAV genotypes, containing a genome of approximately 35 kb (Kurokawa et al (1978) *J. Virol.* 28:212-218). Reddy et al. (1998, *Journal of Virology*, 72:1394) disclose nucleotide sequence, genome organization, and transcription map of BAV3. Reddy et al. (1999, *Journal of Virology*, 73: 9137) disclose a replication-defective BAV3 as an expression vector. BAV3, a representative of subgroup 1 of BAVs (Bartha (1969) *Acta Vet. Acad. Sci. Hung.* 19:319-321), is a common pathogen of cattle usually

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resulting in subclinical infection (Darbyshire *et al.* (1965). *J. Comp. Pathol.* 75:327-330), though occasionally associated with a more serious respiratory tract infection (Darbyshire *et al.*, 1966 *Res. Vet. Sci.* 7:81-93; Mattson *et al.*, 1988 *J. Vet Res* 49:67-69). Like other adenoviruses, BAV3 is a non-enveloped icosahedral particle of 75 nm in diameter (Niiyama *et al.* (1975) *J. Virol.* 16:621-633) containing a linear double-stranded DNA molecule. BAV3 can produce tumors when injected into hamsters (Darbyshire, 1966 *Nature* 211:102) and viral DNA can efficiently effect morphological transformation of mouse, hamster or rat cells in culture (Tsukamoto and Sugino, 1972 *J. Virol.* 9:465-473; Motoi *et al.*, 1972 *Gann* 63:415-418). Cross hybridization was observed between BAV3 and human adenovirus type 2 (HAD2) (Hu *et al.*, 1984 *J. Virol.* 49:604-608) in most regions of the genome including some regions near but not at the left end of the genome.

Porcine adenovirus (PAV) infection has been associated with encephalitis, pneumonia, kidney lesions and diarrhea. See Derbyshire (1992) In: "Diseases of Swine" (ed. Leman *et al.*), 7th edition, Iowa State University Press, Ames, IA. pp. 225-227. It has been shown that PAV is capable of stimulating both humoral response and a mucosal antibody responses in the intestine of infected piglets. Tuboly *et al.* (1993) *Res. in Vet. Sci.* 54:345-350. Cross-neutralization studies have indicated the existence of at least five serotypes of PAV. See Derbyshire *et al.* (1975) *J. Comp. Pathol.* 85:437-443; and Hirahara *et al.* (1990) *Jpn. J. Vet. Sci.* 52:407-409. Previous studies of the PAV genome have included the determination of restriction maps for PAV Type 3 (PAV-3) and cloning of restriction fragments representing the complete genome of PAV-3. See Reddy *et al.* (1993) *Intervirology* 36:161-168. In addition, restriction maps for PAV-1 and PAV-2 have been determined. See Reddy *et al.* (1995b) *Arch. Virol.* 140:195-200.

Nucleotide sequences have been determined for segments of the genome of various PAV serotypes. Sequences of the E3, pVIII and fiber genes of PAV-3 were determined by Reddy *et al.* (1995) *Virus Res.* 36:97-106. The E3, pVIII and fiber

genes of PAV-1 and PAV-2 were sequenced by Reddy *et al.* (1996) *Virus Res.* 43:99-109, while the PAV-4 E3, pVIII and fiber gene sequences were determined by Kleiboeker (1994) *Virus Res.* 31:17-25. The PAV-4 fiber gene sequence was determined by Kleiboeker (1995) *Virus Res.* 39:299-309. Inverted terminal repeat (ITR) sequences for all five PAV serotypes (PAV-1 through PAV-5) were determined by Reddy *et al.* (1995) *Virology* 212:237-239. The PAV-3 penton sequence was determined by McCoy *et al.* (1996) *Arch. Virol.* 141:1367-1375. The nucleotide sequence of the E1 region of PAV-4 was determined by Kleiboeker (1995) *Virus Res.* 36:259-268. The sequence of the protease (23K) gene of PAV-3 was determined by McCoy *et al.* (1996) *DNA Seq.* 6:251-254. The sequence of the PAV-3 hexon gene (and the 14 N-terminal codons of the 23K protease gene) has been deposited in the GenBank database under accession No. U34592. The sequence of the PAV-3 100K gene has been deposited in the GenBank database under accession No. U82628. The sequence of the PAV-3 E4 region has been determined by Reddy *et al.* (1997) *Virus Genes* 15:87-90. Vrati *et al.* (1995, *Virology*, 209:400-408) disclose sequences for ovine adenovirus.

At least 47 serotypes of human adenoviruses have been described. Reviews of the most common serotypes associated with particular diseases have been published. See for example, Foy H.M. (1989) *Adenoviruses* In Evans AS (ed). Viral Infections of Humans. New York, Plenum Publishing, pp 77-89 and Rubin B.A. (1993) *Clinical picture and epidemiology of adenovirus infections*, *Acta Microbiol. Hung* 40:303-323. The capsid of a human adenovirus demonstrates icosahedral symmetry and contains 252 capsomers. The capsomers consist of 240 hexons and 12 pentons with a projecting fiber on each of the pentons. The pentons and hexons are each derived from different viral polypeptides. The fibers, which are responsible for type-specific antibodies, vary in length among human strains. The hexons are group specific complement-fixing antibodies, whereas the pentons are especially active in hemagglutination (Plotkin and Orenstein, Vaccines, 3rd edition, W.B. Saunders

Company Philadelphia, pp609-623). The fiber region assumes a homotrimeric conformation which is necessary for association of the mature fiber protein with the penton base in the formation of the adenovirus capsid. Fiber associates with penton base by virtue of non-covalent interactions between the amino terminus of the fiber trimer and a conserved domain within the penton base. It has been shown that the globular carboxyterminal knob domain of the adenovirus fiber protein is the ligand for attachment to the adenovirus primary cellular receptor (Krasnykh et al. (1996) *Journal of Virology*, 70:6839.). The distal, C-terminal domain of the trimeric fiber molecule terminates in a knob which binds with high affinity to a specific primary receptor. After binding, Arg-Gly-Asp (RGD) motifs in the penton base interact with cellular integrins of the $\alpha\beta 3$ and $\alpha\beta 5$ types which function as secondary receptors. This interaction triggers cellular internalization whereby the virion resides within the endosome. The endosome membrane is lysed in a process mediated by the penton base, releasing the contents of the endosome to the cytoplasm. During these processes, the virion is gradually uncoated and the adenovirus DNA is transported into the nucleus (Shayakhmetov *et al.* (2000) *Journal of Virology* 74:2567-2583).

For general background references regarding adenovirus and development of adenoviral vector systems, see Graham *et al.* (1973) *Virology* 52:456-467; Takiff *et al.* (1981) *Lancet* 11:832-834; Berkner *et al.* (1983) *Nucleic Acid Research* 11: 6003-6020; Graham (1984) *EMBO J* 3:2917-2922; Bett *et al.* (1993) *J. Virology* 67:5911-5921; and Bett *et al.* (1994) *Proc. Natl. Acad. Sci. USA* 91:8802-8806.

Adenoviruses generally undergo a lytic replication cycle following infection of a host cell. In addition to lysing the infected cell, the replicative process of adenovirus blocks the transport and translation host cell mRNA, thus inhibiting cellular protein synthesis. For a review of adenoviruses and adenovirus replication, see Shenk, T. and Horwitz, M.S., *Virology*, third edition, Fields, B.N. *et al.*, eds., Raven Press Limited, New York (1996), Chapters 67 and 68, respectively.

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The application of genetic engineering has resulted in several attempts to prepare adenovirus expression systems for obtaining vaccines. Examples of such research include the disclosures in U.S. Patent 4,510,245 of an adenovirus major late promoter for expression in a yeast host; U.S. Patent 4,920,209 on a live recombinant adenovirus type 7 with a gene coding for hepatitis-B surface antigen located at a deleted early region 3; European Patent 389 286 on a non-defective human adenovirus 5 recombinant expression system in human cells for HCMV major envelope glycoprotein; WO 91/11525 on live non-pathogenic immunogenic viable canine adenovirus in a cell expressing E1A proteins; and French Patent 2 642 767 on vectors containing a leader and/or promoter from the E3 region of adenovirus 2. United States Patent Numbers 6,001,591 and 5,820,868 and International Publication Number WO 95/16048 disclose recombinant protein production in bovine adenovirus expression vector systems. United States Patent Number 5,922,576 discloses systems for generating recombinant adenoviruses.

Krasnykh *et al.* (1996, *Journal of Virology*, 70:6839), Zabner *et al.* (1999) *Journal of Virology*, 73:8689), and Shayakhmetov *et al. supra* report generation of human adenovirus vectors with modified fiber regions. Xu *et al.* (1998, *Virology*, 248:156-163) disclose an ovine adenovirus carrying the fiber protein cell binding domain of human Adenovirus Type 5.

The disclosure of all patents and publications cited herein are incorporated by reference in their entirety.

DISCLOSURE OF THE INVENTION

The present invention provides adenoviruses, preferably bovine adenoviruses, comprising a modification in a polynucleotide encoding a capsid protein, or fragment thereof, wherein said protein, or fragment thereof, is associated with tropism and wherein said modification is associated with altered tropism. The present invention further provides host cells and methods comprising the modified adenoviruses.

Accordingly, the present invention provides bovine adenovirus vectors comprising a modification in a polynucleotide encoding a capsid protein, or fragment thereof, wherein said protein, or fragment thereof, is associated with tropism and wherein said modification is associated with altered tropism. In some embodiments, the

5 polynucleotide encoding a capsid protein, or fragment thereof, is replaced with a polynucleotide encoding a heterologous mammalian capsid protein, or fragment thereof. The capsid protein, or fragment thereof, includes adenovirus penton, hexon or fiber proteins, or fragments thereof. In some embodiments, the modification is in a polynucleotide encoding the knob region of a fiber protein. In other embodiments, a
10 polynucleotide encoding a bovine adenovirus penton, hexon and/or fiber protein(s) is replaced with at least one polynucleotide encoding a heterologous mammalian adenovirus penton, hexon and/or fiber protein(s), respectively. In additional embodiments, a polynucleotide encoding a bovine adenovirus penton protein, or fragment thereof, is replaced with at least one polynucleotide encoding a
15 heterologous mammalian adenovirus penton protein, or fragment thereof; a polynucleotide encoding a bovine adenovirus hexon protein, or fragment thereof, is replaced with at least one polynucleotide encoding a heterologous mammalian adenovirus hexon protein, or fragment thereof; or a polynucleotide encoding a bovine adenovirus fiber protein, or fragment thereof, such as a knob region, is replaced with
20 at least one polynucleotide encoding a heterologous mammalian adenovirus fiber protein, or fragment thereof, such as a heterologous knob region of a fiber protein.

In further embodiments, heterologous mammalian adenoviruses include bovine, porcine, ovine, canine or human adenovirus. In additional embodiments, bovine adenoviruses include sub-type 1 adenovirus, and in particular BAV3, or sub-type 2
25 adenovirus. In other embodiments, the bovine adenovirus vector further comprises a polynucleotide encoding a heterologous protein. In some embodiments, the heterologous protein is a therapeutic protein. In other embodiments, the heterologous protein includes cytokines; lymphokines; membrane receptors recognized by

pathogenic organisms, dystrophins; insulin; proteins participating in cellular ion channels; antisense RNAs; proteins capable of inhibiting the activity of a protein produced by a pathogenic gene, a protein inhibiting an enzyme activity, protein variants of pathogenic proteins; antigenic epitopes; major histocompatibility complex classes I and II proteins; antibodies; immunotoxins; toxins; growth factors or growth hormones; cell receptors or their ligands; tumor suppressors; cellular enzymes; or suicide genes. In yet other embodiments, an adenovirus vector lacks E1 function. In additional embodiments, an adenovirus vector has a deletion in part or all of the E1 gene region. In further embodiments, the adenovirus vector has a deletion of part or all of the E3 gene region. In yet further embodiments, a polynucleotide encoding a heterologous protein is inserted in the adenovirus E1 gene region. In other embodiments, a polynucleotide encoding a heterologous protein is inserted in the adenovirus E3 gene region. In further embodiments, an adenovirus vector is replication-defective, and in yet further embodiments, an adenovirus vector is replication-competent. The present invention also encompasses host cells comprising a bovine adenovirus vector having a modification in a polynucleotide encoding a capsid protein, or fragment thereof.

The present invention also provides methods of producing a recombinant bovine adenovirus vector comprising a modification in a polynucleotide encoding a capsid protein, or fragment thereof, comprising the steps of, obtaining a bovine adenovirus vector; and introducing a modification into a polynucleotide encoding a capsid protein, or fragment thereof, wherein said capsid protein, or fragment thereof, is associated with tropism and wherein said modification is associated with altered tropism. In some embodiments, the modification is a replacement of at least one polynucleotide encoding a bovine adenovirus penton, hexon and/or fiber protein, or fragment thereof, with a heterologous mammalian penton, hexon and/or fiber protein, or fragment thereof. In other embodiments, the modification is a replacement of a polynucleotide encoding a knob region of a fiber protein. In further embodiments,

the adenovirus vector further comprises a polynucleotide encoding a heterologous protein.

The present invention further provides recombinant bovine adenoviruses comprising a modification in a capsid protein, or fragment thereof, wherein said capsid protein, or fragment thereof, is associated with tropism and wherein said modification is associated with altered tropism. In further embodiments, recombinant adenoviruses comprise polynucleotides encoding a heterologous protein. In further embodiments, a polynucleotide encoding a heterologous protein is inserted in the adenovirus E1 gene region; in yet further embodiments, a polynucleotide encoding a heterologous protein is inserted in the adenovirus E3 gene region. In some embodiments, a recombinant adenovirus is replication-competent and in other embodiments, a recombinant adenovirus is replication-defective. In some embodiments, a recombinant adenovirus comprises a replacement of at least one polynucleotide encoding a bovine adenovirus penton, hexon and/or fiber protein(s), or fragment thereof, with a heterologous mammalian penton, hexon and/or fiber protein(s), or fragment thereof. In yet further embodiments, a recombinant adenovirus comprises a modification in a knob region of a fiber protein.

The present invention also provides immunogenic compositions comprising a bovine adenovirus wherein said adenovirus comprises a polynucleotide encoding a modification in a capsid protein, or fragment thereof, and wherein said protein, or fragment thereof, is associated with tropism and wherein said modification is associated with altered tropism. In some embodiments, the capsid protein, or fragment thereof, includes penton, hexon or fiber protein(s), or a fragment thereof, of an adenovirus. In some embodiments of immunogenic compositions, the modification comprises a replacement of a polynucleotide encoding a bovine capsid protein, or fragment thereof, with a polynucleotide encoding a heterologous mammalian adenovirus capsid protein, or fragment thereof. In other embodiments of immunogenic compositions, the modification comprises a replacement of a

polynucleotide encoding a bovine knob region of a fiber protein with a
polynucleotide encoding a heterologous mammalian adenovirus knob region of a
fiber protein. In other embodiments, the bovine adenovirus is a sub-type 1
adenovirus, in particular, BAV3, or a sub-type 2 adenovirus. In additional
5 embodiments, immunogenic compositions comprise a bovine adenovirus comprises a
polynucleotide encoding a heterologous protein. In other embodiments,
immunogenic compositions comprise a bovine adenovirus comprising a
polynucleotide encoding cytokines; lymphokines; membrane receptors recognized by
pathogenic organisms, dystrophins; insulin; proteins participating in cellular ion
10 channels; antisense RNAs; proteins capable of inhibiting the activity of a protein
produced by a pathogenic gene, a protein inhibiting an enzyme activity, protein
variants of pathogenic proteins; antigenic epitopes; major histocompatibility complex
classes I and II proteins; antibodies; immunotoxins; toxins; growth factors or growth
hormones; cell receptors or their ligands; tumor suppressors; cellular enzymes; or
15 suicide genes.

 The present invention also encompasses pharmaceutical compositions capable
of inducing an immune response in a mammalian subject. In some embodiments,
pharmaceutical compositions comprise an immunogenic composition comprising a
bovine adenovirus having a modified capsid protein, or fragment thereof, wherein the
20 protein, or fragment thereof, is associated with tropism and wherein the modification
is associated with altered tropism. In some embodiments of the pharmaceutical
compositions, immunogenic compositions comprise bovine adenovirus vectors
comprising a polynucleotide encoding a heterologous protein. In some embodiments,
the heterologous protein is a therapeutic protein. In other embodiments, the
25 pharmaceutical composition further comprises a pharmaceutically acceptable
excipient.

 The present invention also provides methods for eliciting an immune response
in a mammalian host to protect against infection, the method comprising

administering a pharmaceutical composition of the present invention to a mammalian host in need. The present invention also provides methods of gene delivery in a mammalian host, the methods comprising administering to the host a bovine adenovirus vector comprising a polynucleotide encoding a modified capsid protein, or fragment thereof, wherein the protein is associated with tropism and wherein the modification is associated with altered tropism and wherein the adenovirus vector further comprises a polynucleotide encoding a heterologous protein. In some embodiments, the heterologous polynucleotide encodes a therapeutic protein.

BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1A-1S shows the complete nucleotide sequence of the BAV3 genome. In the polynucleotide sequence for BAV3, the penton regions starts at 12919 and ends at 14367; the hexon region starts at 17809 and ends at 20517; the fiber region starts at 27968 and ends at 30898. The knob domain of the fiber region starts after the 4 residues, TLWT, as shown in Figure 4.

Figure 2 shows a transcriptional map of the BAV3 genome, derived from transcriptional mapping of mRNAs and sequencing of cDNA clones.

Figure 3 illustrates the construction of BAV600 that expresses the HAV-5 fiber knob protein.

Figure 4 illustrates the characterization of BAV600.

Figures 5A-5B shows the analysis of BAV600 by Restriction Enzyme *Bgl*II digestion. Figure 5A depicts a gel electrophoresis and Figure 5B depicts a Southern Blot.

Figure 6 shows the expression of HAV-5 fiber Knob by BAV600.

Figures 7A-7B show the transduction of Human cell lines by BAV600. Figure 7A show results of an MOI of 1 whereas Figure 7B shows results of an MOI of 5.

Figure 8 shows a FACS analysis of BAV304 and BAV600 transduction of Human cells.

Figure 9 shows the expression of early and late BAV-3 proteins in human cell lines, HeLa, HEp-2, A549, 293 and MDBK.

5 Figure 10 illustrates BAV3 replication in human cells.

Figure 11 shows the neutralization of BAV600 by a monoclonal antibody specific for HAV-5 fiber knob region.

Ans A3
~~Figure 12 depicts the amino acid sequence for Human adenovirus 5 (HAV-5) fiber protein.~~

Ans A4
~~Figure 13 depicts the amino acid sequence for the Bovine Adenovirus 3 (BAV-3) fiber protein.~~

Ans A5
~~Figure 14 depicts the amino acid sequence of Ovine Adenovirus 287 fiber protein.~~

Ans A6
~~Figure 15 shows the amino acid sequence of Porcine Adenovirus 3 (PAV-3) fiber protein.~~

Ans A7
~~Figure 16 shows the amino acid sequence of Canine Adenovirus 2 (CAV-2) fiber protein.~~

Ans A8
~~Figures 17A-17G depicts an amino acid alignment of various mammalian adenovirus fiber regions using the clustal method of the Multialign program.~~

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BEST MODE FOR CARRYING OUT THE INVENTION

We have discovered and constructed improved adenovirus vectors, in particular improved bovine adenovirus vectors, having altered tropism. The bovine adenovirus vectors of the present invention comprise a modification in a polynucleotide encoding at least one capsid protein, wherein the protein, or fragment thereof, is associated with tropism and wherein the modification is associated with altered tropism.

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Capsid proteins include penton, hexon and fiber proteins. In one embodiment illustrated herein, a BAV3 adenovirus vector was constructed, BAV600, which comprised a replacement of the BAV3 fiber knob region with a human adenovirus (Ad5) fiber knob region. BAV600 demonstrated increased transduction in human cell lines as compared to a control adenovirus.

The present invention encompasses bovine adenovirus vectors comprising a replacement of a capsid protein, or fragment thereof, with a heterologous mammalian capsid protein, or fragment thereof, as long as the protein is associated with tropism and the replacement is associated with altered tropism. For example, in one embodiment, a bovine knob domain of a fiber protein is replaced with a porcine or ovine knob region of a fiber protein in order to alter species tropism. Such a bovine adenovirus vector can be used as an immunogen to boost immunity in a porcine or ovine mammal that has been primed with a porcine or ovine adenovirus, respectively. In such an immunization protocol, a boost immunization is achieved by administration of the bovine adenovirus having species specificity for the porcine or ovine mammal, while avoiding the affect of any neutralizing antibodies against the porcine or ovine mammal produced as a result of the priming immunization.

Alternatively, in another embodiment, a bovine fiber protein, or fragment thereof, such as the knob region, is replaced with a heterologous bovine fiber protein, or fragment thereof, such as a knob region of a fiber protein in order to alter bovine cell specificity. For one example, a bovine adenovirus sub-type 1 fiber region, or fragment thereof, such as a knob domain, is replaced with a bovine adenovirus sub-type 2 fiber region, or fragment thereof, such as a knob domain, in order to alter bovine cell-type specificity. Such a bovine adenovirus vector can be used as an immunogen to target specific cells or tissues.

The invention also encompasses the use of a bovine adenovirus comprising a replacement of a bovine capsid protein, or fragment thereof, with a human adenovirus capsid protein, or fragment thereof, such that the modified bovine adenovirus has

species specificity for humans. Such bovine adenoviruses can be used in human immunization protocols, where preexisting neutralizing antibodies against human adenovirus -5 (HAV-5) in clinical patients may present an obstacle for efficient use of HAV-5.

5 Additionally, to provide a therapeutic effect to target cells, one or more heterologous therapeutic proteins may be present in the adenovirus vector.

Definitions

In describing the present invention, the following terminology, as defined below, will be used.

10 An "adenovirus vector" or "adenoviral vector" (used interchangeably) comprises a polynucleotide construct of the invention. A polynucleotide construct of this invention may be in any of several forms, including, but not limited to, DNA, DNA encapsulated in an adenovirus coat, DNA packaged in another viral or viral-like form (such as herpes simplex, and AAV), DNA encapsulated in liposomes, DNA
15 complexed with polylysine, complexed with synthetic polycationic molecules, conjugated with transferrin, and complexed with compounds such as PEG to immunologically "mask" the molecule and/or increase half-life, and conjugated to a nonviral protein. Preferably, the polynucleotide is DNA. As used herein, "DNA" includes not only bases A, T, C, and G, but also includes any of their analogs or
20 modified forms of these bases, such as methylated nucleotides, internucleotide modifications such as uncharged linkages and thioates, use of sugar analogs, and modified and/or alternative backbone structures, such as polyamides. Adenovirus vectors may be replication-competent or replication-defective in a target cell.

 As used herein, the term "altered tropism" refers to changing the specificity of an
25 adenovirus. The term "altered tropism" encompasses changing species specificity as well as changing tissue or cell specificity of an adenovirus. In embodiments illustrated herein, species specificity is altered by producing modifications in a capsid

protein(s), or fragment thereof, such as the fiber protein, and in particular the knob region of a fiber protein.

A "capsid protein" as used herein includes penton, hexon and fiber regions of an adenovirus. A capsid protein is associated with tropism if it directly or indirectly affects adenovirus tropism. A "modification of a capsid protein associated with altered tropism" as used herein refers to producing an alteration of a polynucleotide encoding a capsid protein, ie, a penton, hexon or fiber protein region, or fragment thereof, such as the knob domain of the fiber region such that specificity is altered.

"Associated with" means that the modification contributes to the altered tropism either directly or indirectly. In embodiments illustrated herein, the modification is a replacement of bovine capsid protein regions with a heterologous mammalian capsid protein region in order to produce species specificity in the adenovirus. Replacement of one species capsid protein region with a heterologous capsid protein region may also produce altered tissue or cell specificity.

A "replicon" is any genetic element (e.g., plasmid, chromosome, virus) that functions as an autonomous unit of DNA replication *in vivo*; i.e., is capable of replication under its own control.

As used herein, the term "vector" refers to a polynucleotide construct designed for transduction/transfection of one or more cell types. Vectors may be, for example, "cloning vectors" which are designed for isolation, propagation and replication of inserted nucleotides, "expression vectors" which are designed for expression of a nucleotide sequence in a host cell, or a "viral vector" which is designed to result in the production of a recombinant virus or virus-like particle, or "shuttle vectors", which comprise the attributes of more than one type of vector.

By "live virus" is meant, in contradistinction to "killed" virus, a virus which is capable of producing identical progeny in tissue culture and inoculated animals.

A "helper-free virus vector" is a vector that does not require a second virus or a cell line to supply something defective in the vector.

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A "double-stranded DNA molecule" refers to the polymeric form of deoxyribonucleotides (adenine, guanine, thymine, or cytosine) in its normal, double-stranded helix. This term refers only to the primary and secondary structure of the molecule, and does not limit it to any particular tertiary forms. Thus, this term includes double-stranded DNA found, inter alia, in linear DNA molecules (e.g., restriction fragments of DNA from viruses, plasmids, and chromosomes). In discussing the structure of particular double-stranded DNA molecules, sequences may be described herein according to the normal convention of giving only the sequence in the 5' to 3' direction along the nontranscribed strand of DNA (i.e., the strand having the sequence homologous to the mRNA).

A DNA "coding sequence" is a DNA sequence which is transcribed and translated into a polypeptide *in vivo* when placed under the control of appropriate regulatory sequences. The boundaries of the coding sequence are determined by a start codon at the 5' (amino) terminus and a translation stop codon at the 3' (carboxy) terminus. A coding sequence can include, but is not limited to, procaryotic sequences, cDNA from eucaryotic mRNA, genomic DNA sequences from eucaryotic (e.g., mammalian) DNA, viral DNA, and even synthetic DNA sequences. A polyadenylation signal and transcription termination sequence will usually be located 3' to the coding sequence.

A "transcriptional promoter sequence" is a DNA regulatory region capable of binding RNA polymerase in a cell and initiating transcription of a downstream (3' direction) coding sequence. For purposes of defining the present invention, the promoter sequence is bound at the 3' terminus by the translation start codon (ATG) of a coding sequence and extends upstream (5' direction) to include the minimum number of bases or elements necessary to initiate transcription at levels detectable above background. Within the promoter sequence will be found a transcription initiation site (conveniently defined by mapping with nuclease S1), as well as protein binding domains (consensus sequences) responsible for the binding of RNA

polymerase. Eucaryotic promoters will often, but not always, contain "TATA" boxes and "CAAT" boxes. Procaryotic promoters contain Shine-Dalgarno sequences in addition to the -10 and -35 consensus sequences.

5 DNA "control sequences" refer collectively to promoter sequences, ribosome binding sites, splicing signals, polyadenylation signals, transcription termination sequences, upstream regulatory domains, enhancers, translational termination sequences and the like, which collectively provide for the transcription and translation of a coding sequence in a host cell.

10 A coding sequence or sequence encoding a protein is "operably linked to" or "under the control of" control sequences in a cell when RNA polymerase will bind the promoter sequence and transcribe the coding sequence into mRNA, which is then translated into the polypeptide encoded by the coding sequence.

A "host cell" is a cell which has been transformed, or is capable of transformation, by an exogenous DNA sequence.

15 A cell has been "transformed" by exogenous DNA when such exogenous DNA has been introduced inside the cell membrane. Exogenous DNA may or may not be integrated (covalently linked) to chromosomal DNA making up the genome of the cell. In procaryotes and yeasts, for example, the exogenous DNA may be maintained on an episomal element, such as a plasmid. A stably transformed cell is one in which the exogenous DNA has become integrated into the chromosome so that it is inherited by daughter cells through chromosome replication. For mammalian cells, this stability is demonstrated by the ability of the cell to establish cell lines or clones comprised of a population of daughter cell containing the exogenous DNA.

20 A "clone" is a population of daughter cells derived from a single cell or common ancestor. A "cell line" is a clone of a primary cell that is capable of stable growth *in vitro* for many generations.

25 A "heterologous" region of a DNA construct is an identifiable segment of DNA within or attached to another DNA molecule that is not found in association

with the other molecule in nature. Thus, when the heterologous region encodes a viral gene, the gene will usually be flanked by DNA that does not flank the viral gene in the genome of the source virus or virus-infected cells. Another example of the heterologous coding sequence is a construct wherein the coding sequence itself is not found in nature (e.g., synthetic sequences having codons different from the native gene). Allelic variation or naturally occurring mutational events do not give rise to a heterologous region of DNA, as used herein. As used herein in describing adenovirus vectors, "heterologous mammalian capsid region" means that the capsid region is obtainable from another mammalian species of adenovirus or is obtainable from the same species mammal but from a different type or sub-type adenovirus. For example "heterologous mammalian capsid protein" encompasses replacement of one sub-type bovine adenovirus capsid protein with another sub-type bovine adenovirus capsid protein as well as replacement of a bovine adenovirus capsid protein with another species capsid protein, such as a human capsid protein, as well as replacement of bovine adenovirus capsid proteins regions with another serotype bovine adenovirus capsid protein.

"Bovine host" refers to cattle of any breed, adult or infant.

The term "protein" is used herein to designate a polypeptide or glycosylated polypeptide, respectively, unless otherwise noted. The term "polypeptide" is used in its broadest sense, i.e., any polymer of amino acids (dipeptide or greater) linked through peptide bonds. Thus, the term "polypeptide" includes proteins, oligopeptides, protein fragments, analogs, muteins, fusion proteins and the like.

"Native" proteins or polypeptides refer to proteins or polypeptides recovered from adenovirus or adenovirus-infected cells. Thus, the term "native BAV polypeptide" would include naturally occurring BAV proteins and fragments thereof. "Non-native" polypeptides refer to polypeptides that have been produced by recombinant DNA methods or by direct synthesis. "Recombinant" polypeptides

refers to polypeptides produced by recombinant DNA techniques; i.e., produced from cells transformed by an exogenous DNA construct encoding the desired polypeptide.

A "substantially pure" protein will be free of other proteins, preferably at least 10% homogeneous, more preferably 60% homogeneous, and most preferably 95% homogeneous.

An "antigen" refers to a molecule containing one or more epitopes that will stimulate a host's immune system to make a humoral and/or cellular antigen-specific response. The term is also used interchangeably with "immunogen."

A "hapten" is a molecule containing one or more epitopes that does not stimulate a host's immune system to make a humoral or cellular response unless linked to a carrier.

The term "epitope" refers to the site on an antigen or hapten to which a specific antibody molecule binds or is recognized by T cells. The term is also used interchangeably with "antigenic determinant" or "antigenic determinant site."

An "immunological response" to a composition or vaccine is the development in the host of a cellular and/or antibody-mediated immune response to the composition or vaccine of interest. Usually, such a response consists of the subject producing antibodies, B cells, helper T cells, suppressor T cells, and/or cytotoxic T cells directed specifically to an antigen or antigens included in the composition or vaccine of interest.

The terms "immunogenic polypeptide" and "immunogenic amino acid sequence" and "immunogen" refer to a polypeptide or amino acid sequence, respectively, which elicit antibodies that neutralize viral infectivity, and/or mediate antibody-complement or antibody-dependent cell cytotoxicity to provide protection of an immunized host. An "immunogenic polypeptide" as used herein, includes the full length (or near full length) sequence of the desired protein or an immunogenic fragment thereof.

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By "immunogenic fragment" is meant a fragment of a polypeptide which includes one or more epitopes and thus elicits antibodies that neutralize viral infectivity, and/or mediates antibody-complement or antibody-dependent cell cytotoxicity to provide protection of an immunized host. Such fragments will usually be at least about 5 amino acids in length, and preferably at least about 10 to 15 amino acids in length. There is no critical upper limit to the length of the fragment, which could comprise nearly the full length of the protein sequence, or even a fusion protein comprising fragments of two or more of the antigens. The term "treatment" as used herein refers to treatment of a mammal, such as bovine or human or other mammal, either (i) the prevention of infection or reinfection (prophylaxis), or (ii) the reduction or elimination of symptoms of an infection. The vaccine comprises the recombinant BAV itself or recombinant antigen produced by recombinant BAV.

By "infectious" is meant having the capacity to deliver the viral genome into cells.

The terms "polynucleotide" and "nucleic acid", used interchangeably herein, refer to a polymeric form of nucleotides of any length, either ribonucleotides or deoxyribonucleotides. These terms include a single-, double- or triple-stranded DNA, genomic DNA, cDNA, RNA, DNA-RNA hybrid, or a polymer comprising purine and pyrimidine bases, or other natural, chemically, biochemically modified, non-natural or derivatized nucleotide bases. The backbone of the polynucleotide can comprise sugars and phosphate groups (as may typically be found in RNA or DNA), or modified or substituted sugar or phosphate groups. Alternatively, the backbone of the polynucleotide can comprise a polymer of synthetic subunits such as phosphoramidates and thus can be a oligodeoxynucleoside phosphoramidate (P-NH₂) or a mixed phosphoramidate- phosphodiester oligomer. Peyrottes et al. (1996) *Nucleic Acids Res.* 24: 1841-8; Chaturvedi et al. (1996) *Nucleic Acids Res.* 24: 2318-23; Schultz et al. (1996) *Nucleic Acids Res.* 24: 2966-73. A phosphorothioate linkage can be used in place of a phosphodiester linkage. Braun et al. (1988) *J. Immunol.* 141:

2084-9; Latimer et al. (1995) *Molec. Immunol.* 32: 1057-1064. In addition, a double-stranded polynucleotide can be obtained from the single stranded polynucleotide product of chemical synthesis either by synthesizing the complementary strand and annealing the strands under appropriate conditions, or by synthesizing the complementary strand de novo using a DNA polymerase with an appropriate primer. Reference to a polynucleotide sequence (such as referring to a SEQ ID NO) also includes the complement sequence.

The following are non-limiting examples of polynucleotides: a gene or gene fragment, exons, introns, mRNA, tRNA, rRNA, ribozymes, cDNA, recombinant polynucleotides, branched polynucleotides, plasmids, vectors, isolated DNA of any sequence, isolated RNA of any sequence, nucleic acid probes, and primers. A polynucleotide may comprise modified nucleotides, such as methylated nucleotides and nucleotide analogs, uracyl, other sugars and linking groups such as fluororibose and thioate, and nucleotide branches. The sequence of nucleotides may be interrupted by non-nucleotide components. A polynucleotide may be further modified after polymerization, such as by conjugation with a labeling component. Other types of modifications included in this definition are caps, substitution of one or more of the naturally occurring nucleotides with an analog, and introduction of means for attaching the polynucleotide to proteins, metal ions, labeling components, other polynucleotides, or a solid support. Preferably, the polynucleotide is DNA. As used herein, "DNA" includes not only bases A, T, C, and G, but also includes any of their analogs or modified forms of these bases, such as methylated nucleotides, internucleotide modifications such as uncharged linkages and thioates, use of sugar analogs, and modified and/or alternative backbone structures, such as polyamides.

A polynucleotide or polynucleotide region has a certain percentage (for example, 80%, 85%, 90%, or 95%) of "sequence identity" to another sequence means that, when aligned, that percentage of bases are the same in comparing the two sequences. This alignment and the percent homology or sequence identity can be

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determined using software programs known in the art, for example those described in *Current Protocols in Molecular Biology* (F.M. Ausubel et al., eds., 1987) Supplement 30, section 7.7.18, Table 7.7.1. A preferred alignment program is ALIGN Plus (Scientific and Educational Software, Pennsylvania), preferably using
5 default parameters, which are as follows: mismatch = 2; open gap = 0; extend gap = 2.

“Under transcriptional control” is a term well understood in the art and indicates that transcription of a polynucleotide sequence, usually a DNA sequence, depends on its being operably (operatively) linked to an element which contributes to
10 the initiation of, or promotes, transcription. “Operably linked” refers to a juxtaposition wherein the elements are in an arrangement allowing them to function.

As used herein, “cytotoxicity” is a term well understood in the art and refers to a state in which a cell’s usual biochemical or biological activities are compromised (i.e., inhibited). These activities include, but are not limited to, metabolism; cellular
15 replication; DNA replication; transcription; translation; uptake of molecules. “Cytotoxicity” includes cell death and/or cytolysis. Assays are known in the art which indicate cytotoxicity, such as dye exclusion, ³H-thymidine uptake, and plaque assays.

In the context of adenovirus, a “heterologous polynucleotide” or
20 “heterologous gene” or “transgene” is any polynucleotide or gene that is not present in wild-type adenovirus. Preferably, the transgene will also not be expressed or present in the target cell prior to introduction by the adenovirus vector.

In the context of adenovirus, a “heterologous” promoter or enhancer is one which is not associated with or derived from an adenovirus gene.

25 In the context of adenovirus, an “endogenous” promoter, enhancer, or control region is native to or derived from adenovirus.

A “host cell” includes an individual cell or cell culture which can be or has been a recipient of an adenoviral vector(s) of this invention. Host cells include

progeny of a single host cell, and the progeny may not necessarily be completely identical (in morphology or in total DNA complement) to the original parent cell due to natural, accidental, or deliberate mutation and/or change. A host cell includes cells transfected or infected *in vivo* or *in vitro* with an adenoviral vector of this invention.

5 “Replication” and “propagation” are used interchangeably and refer to the ability of an adenovirus vector of the invention to reproduce or proliferate. These terms are well understood in the art. For purposes of this invention, replication involves production of adenovirus proteins and is generally directed to reproduction of adenovirus. Replication can be measured using assays standard in the art and
10 described herein, such as a burst assay or plaque assay. “Replication” and “propagation” include any activity directly or indirectly involved in the process of virus manufacture, including, but not limited to, viral gene expression; production of viral proteins, nucleic acids or other components; packaging of viral components into complete viruses; and cell lysis.

15 A polynucleotide sequence that is “depicted in” a SEQ ID NO means that the sequence is present as an identical contiguous sequence in the SEQ ID NO. The term encompasses portions, or regions of the SEQ ID NO as well as the entire sequence contained within the SEQ ID NO.

20 A “biological sample” encompasses a variety of sample types obtained from an individual and can be used in a diagnostic or monitoring assay. The definition encompasses blood and other liquid samples of biological origin, solid tissue samples such as a biopsy specimen or tissue cultures or cells derived therefrom, and the progeny thereof. The definition also includes samples that have been manipulated in any way after their procurement, such as by treatment with reagents, solubilization, or
25 enrichment for certain components, such as proteins or polynucleotides. The term “biological sample” encompasses a clinical sample, and also includes cells in culture, cell supernatants, cell lysates, serum, plasma, biological fluid, and tissue samples.

An "individual" or "mammalian subject" is a vertebrate, preferably a mammal, more preferably a human. Mammals include, but are not limited to, farm animals, sport animals, rodents, primates, and pets.

An "effective amount" is an amount sufficient to effect beneficial or desired results, including clinical results. An effective amount can be administered in one or more administrations. For purposes of this invention, an effective amount of an adenoviral vector is an amount that is sufficient to palliate, ameliorate, stabilize, reverse, slow or delay the progression of the disease state.

"Expression" includes transcription and/or translation.

As used herein, the term "comprising" and its cognates are used in their inclusive sense; that is, equivalent to the term "including" and its corresponding cognates.

"A," "an" and "the" include plural references unless the context clearly dictates otherwise.

Detailed Description

The present invention identifies capsid proteins associated with tropism and provides methods of constructing adenovirus vectors and recombinant adenoviruses having altered tropism. In preferred embodiments, the adenovirus is a bovine adenovirus, such as a sub-type 1 adenovirus, in particular BAV3, or a sub-type 2 adenovirus. In illustrative embodiments, part or all of a bovine capsid protein encoding polynucleotide sequence associated with tropism is deleted and replaced with part or all of a heterologous mammalian capsid protein encoding polynucleotide sequence which alters adenovirus tropism. In a particular embodiment disclosed herein, the knob region of a bovine fiber protein is replaced with a human knob region of a fiber protein. The present invention also encompasses adenoviruses comprising the replacement of one bovine serotype adenovirus capsid protein associated with tropism with a heterologous bovine serotype adenovirus capsid protein associated with tropism in order to alter cell specificity.

The complete nucleotide sequence of the BAV3 genome is disclosed herein. See Figure 1 (SEQ ID NO 1). A transcriptional map of the BAV3 genome, derived from transcriptional mapping of mRNAs and sequencing of cDNA clones, is presented in Figure 2. Although the size (34,446 bp) and the overall organization of the BAV3 genome appear to be similar to that of HAVs, there are certain differences. Reddy *et al.* (1998) *supra*. One of the distinctive features of the BAV3 genome is the relatively small size of the E3 coding region (1517 bp). Mittal *et al.* (1992) *J. Gen. Virol.* 73:3295-3300; Mittal *et al.* (1993). *J. Gen. Virol.* 74:2825; and Reddy *et al.* (1998) *supra*. Analysis of the sequence of the BAV3 E3 region and its RNA transcripts suggests that BAV3 E3 may encode at least four proteins, one of which (121R) exhibits limited homology with the 14.7 kDa protein of HAV5. Idamakanti (1998) "Molecular characterization of E3 region of bovine adenovirus-3," M.Sc. thesis, University of Saskatchewan, Saskatoon, Saskatchewan.

Reddy *et al.* (1998) *Journal of Virology* 72:1394) disclose nucleotide sequences for BAV3. In the polynucleotide sequence for BAV3, the penton regions starts at 12919 and ends at 14367; the hexon region starts at 17809 and ends at 20517; the fiber region starts at 27968 and ends at 30898. The knob region (or domain) of the fiber protein starts after the residues TLWT motif as shown in Figure 4. The fiber protein also contains shaft and tail regions (or domains).

Human adenoviruses Ad3, Ad4, Ad5, Ad9 and Ad35 are available from the American Tissue Culture Collection ATCC). The National Center for Biotechnology Information GenBank accession number for Ad5 is M73260/M29978; for Ad9 X74659; and for Ad35, U10272. Chow *et al.* (1977, *Cell* 12:1-8) disclose human adenovirus 2 sequences; Davison *et al.* (1993, *J. Mole. Biol.* 234:1308-1316) disclose the DNA sequence of human adenovirus type 40; Sprengel *et al.* (1994, *J. Virol.* 68:379-389) disclose the DNA sequence for human adenovirus type 12 DNA; Vrati *et al.* (1995, *Virology*, 209:400-408) disclose sequences for ovine adenovirus; Morrison *et al.* (1997, *J. Gen. Virol.* 78:873-878) disclose canine adenovirus type 1 DNA

sequence; and Reddy *et al.* (1998, *Virology*, 251:414) disclose DNA sequences for porcine adenovirus.

Shayakhmetov *et al.*, *supra*, provide PCR primers for human Ad9 and human Ad35 fiber regions. The HAV-5 fiber protein is depicted in Figure 12; Figure 13 depicts the amino acid sequence for the Bovine Adenovirus-3 (BAV-3) fiber protein; Figure 14 depicts the amino acid sequence of Ovine Adenovirus 287 fiber protein; Figure 15 depicts the amino acid sequence of Porcine Adenovirus-3 (PAV-3) fiber protein; Figure 16 depicts the amino acid sequence of Canine Adenovirus -2 (CAV-2) fiber protein; and Figures 17A-17G depicts an amino acid alignment of mammalian adenovirus fiber regions using the clustal method of the multialign program. The knob domain of the fiber regions typically starts after the amino acid residue motif TLWT (hinge region), see Figure 4 (one exception is the ovine adenovirus fiber region).

Adenovirus vector constructs can then undergo recombination *in vitro* or *in vivo*, with a BAV genome either before or after transformation or transfection of an appropriate host cell.

Suitable host cells include any cell that will support recombination between a BAV genome and a plasmid containing BAV sequences, or between two or more plasmids, each containing BAV sequences. Recombination is generally performed in procaryotic cells, such as *E. coli*, while transfection of a plasmid containing a viral genome, to generate virus particles, is conducted in eukaryotic cells, preferably mammalian cells, more preferably bovine cell cultures, most preferably MDBK or PFBR cells, and their equivalents. The growth of bacterial cell cultures, as well as culture and maintenance of eukaryotic cells and mammalian cell lines are procedures which are well-known to those of skill in the art.

One or more heterologous polynucleotide sequences can be inserted into one or more regions of the BAV genome to generate a recombinant BAV, limited only by the insertion capacity of the BAV genome and ability of the recombinant BAV to

express the inserted heterologous sequences. In general, adenovirus genomes can accept inserts of approximately 5% of genome length and remain capable of being packaged into virus particles. The insertion capacity can be increased by deletion of non-essential regions and/or deletion of essential regions, such as, for example, E1 function, whose function is provided by a helper cell line, such as one providing E1 function. In some embodiments, a heterologous polynucleotide encoding a protein is inserted into an adenovirus E1 gene region. In some embodiments, an adenovirus has a deletion of part or all of the E1 gene region and is propagated in a helper cell line providing E1 function. In yet other embodiments, a heterologous polynucleotide encoding a protein is inserted into an adenovirus E3 gene region. In other embodiments, an adenovirus has a deletion of part or all of the E3 region.

In one embodiment of the invention, insertion can be achieved by constructing a plasmid containing the region of the BAV genome into which insertion is desired, such as a polynucleotide encoding a capsid protein. Additionally, a polynucleotide encoding a desired therapeutic protein can be inserted into the bovine adenovirus. The plasmid is then digested with a restriction enzyme having a recognition sequence in the BAV portion of the plasmid, and a heterologous polynucleotide sequence is inserted at the site of restriction digestion. The plasmid, containing a portion of the BAV genome with an inserted heterologous sequence, is co-transformed, along with a BAV genome or a linearized plasmid containing a BAV genome, into a bacterial cell (such as, for example, *E. coli*), wherein the BAV genome can be a full-length genome or can contain one or more deletions. Homologous recombination between the plasmids generates a recombinant BAV genome containing inserted heterologous sequences.

Deletion of BAV sequences, to provide a site for insertion of heterologous sequences or to provide additional capacity for insertion at a different site, can be accomplished by methods well-known to those of skill in the art. For example, for BAV sequences cloned in a plasmid, digestion with one or more restriction enzymes

(with at least one recognition sequence in the BAV insert) followed by ligation will, in some cases, result in deletion of sequences between the restriction enzyme recognition sites. Alternatively, digestion at a single restriction enzyme recognition site within the BAV insert, followed by exonuclease treatment, followed by ligation will result in deletion of BAV sequences adjacent to the restriction site. A plasmid containing one or more portions of the BAV genome with one or more deletions, constructed as described above, can be co-transfected into a bacterial cell along with a BAV genome (full-length or deleted) or a plasmid containing either a full-length or a deleted BAV genome to generate, by homologous recombination, a plasmid containing a recombinant BAV genome with a deletion at one or more specific sites. BAV virions containing the deletion can then be obtained by transfection of mammalian cells (including, but not limited to, MDBK or PFBR cells and their equivalents) with the plasmid containing the recombinant BAV genome.

In one embodiment of the invention, insertion sites are adjacent to and downstream (in the transcriptional sense) of BAV promoters. Locations of BAV promoters, and restriction enzyme recognition sequences downstream of BAV promoters, for use as insertion sites, can be easily determined by one of skill in the art from the BAV nucleotide sequence provided herein. Alternatively, various *in vitro* techniques can be used for insertion of a restriction enzyme recognition sequence at a particular site, or for insertion of heterologous sequences at a site that does not contain a restriction enzyme recognition sequence. Such methods include, but are not limited to, oligonucleotide-mediated heteroduplex formation for insertion of one or more restriction enzyme recognition sequences (*see*, for example, Zoller *et al.* (1982) *Nucleic Acids Res.* 10:6487-6500; Brennan *et al.* (1990) *Roux's Arch. Dev. Biol.* 199:89-96; and Kunkel *et al.* (1987) *Meth. Enzymology* 154:367-382) and PCR-mediated methods for insertion of longer sequences. *See*, for example, Zheng *et al.* (1994) *Virus Research* 31:163-186.

It is also possible to obtain expression of a heterologous sequence inserted at a site that is not downstream from a BAV promoter, if the heterologous sequence additionally comprises transcriptional regulatory sequences that are active in eukaryotic cells. Such transcriptional regulatory sequences can include cellular promoters such as, for example, the bovine hsp70 promoter and viral promoters such as, for example, herpesvirus, adenovirus and papovavirus promoters and DNA copies of retroviral long terminal repeat (LTR) sequences.

In another embodiment, homologous recombination in a procaryotic cell can be used to generate a cloned BAV genome; and the cloned BAV genome can be propagated as a plasmid. See for example, U.S. patent 5,922,576. Infectious virus can be obtained by transfection of mammalian cells with the cloned BAV genome rescued from plasmid-containing cells.

The invention also provides BAV regulatory sequences which can be used to regulate the expression of heterologous genes. A regulatory sequence can be, for example, a transcriptional regulatory sequence, a promoter, an enhancer, an upstream regulatory domain, a splicing signal, a polyadenylation signal, a transcriptional termination sequence, a translational regulatory sequence, a ribosome binding site and a translational termination sequence.

In another embodiment, the cloned BAV genome can be propagated as a plasmid and infectious virus can be rescued from plasmid-containing cells.

The presence of viral nucleic acids can be detected by techniques known to one of skill in the art including, but not limited to, hybridization assays, polymerase chain reaction, and other types of amplification reactions. Similarly, methods for detection of proteins are well-known to those of skill in the art and include, but are not limited to, various types of immunoassay, ELISA, Western blotting, enzymatic assay, immunohistochemistry, *etc.* Diagnostic kits comprising the nucleotide sequences of the invention may also contain reagents for cell disruption and nucleic acid purification, as well as buffers and solvents for the formation, selection and

detection of hybrids. Diagnostic kits comprising the polypeptides or amino acid sequences of the invention may also comprise reagents for protein isolation and for the formation, isolation, purification and/or detection of immune complexes.

Various foreign genes or nucleotide sequences or coding sequences
5 (prokaryotic, and eukaryotic) can be inserted in the bovine adenovirus nucleotide sequence, e.g., DNA, in accordance with the present invention, particularly to provide protection against a wide range of diseases and many such genes are already known in the art. The problem heretofore has been to provide a safe, convenient and effective vaccine vector for the genes or sequences, as well as safe, effective means
10 for gene transfer to be used in various gene therapy applications.

An exogenous (*i.e.*, foreign) nucleotide sequence can consist of one or more gene(s) of interest, and preferably of therapeutic interest. In the context of the present invention, a gene of interest can code either for an antisense RNA, a ribozyme or for an mRNA which will then be translated into a protein of interest. A gene of interest
15 can be of genomic type, of complementary DNA (cDNA) type or of mixed type (minigene, in which at least one intron is deleted). It can code for a mature protein, a precursor of a mature protein, in particular a precursor intended to be secreted and accordingly comprising a signal peptide, a chimeric protein originating from the fusion of sequences of diverse origins, or a mutant of a natural protein displaying
20 improved or modified biological properties. Such a mutant may be obtained by, deletion, substitution and/or addition of one or more nucleotide(s) of the gene coding for the natural protein, or any other type of change in the sequence encoding the natural protein, such as, for example, transposition or inversion.

A gene of interest may be placed under the control of elements (DNA control
25 sequences) suitable for its expression in a host cell. Suitable DNA control sequences are understood to mean the set of elements needed for transcription of a gene into RNA (antisense RNA or mRNA) and for the translation of an mRNA into protein. Among the elements needed for transcription, the promoter assumes special

importance. It can be a constitutive promoter or a regulatable promoter, and can be isolated from any gene of eukaryotic, prokaryotic or viral origin, and even adenoviral origin. Alternatively, it can be the natural promoter of the gene of interest. Generally speaking, a promoter used in the present invention may be modified so as to contain regulatory sequences. As examples, a gene of interest in use in the present invention is placed under the control of the promoter of the immunoglobulin genes when it is desired to target its transfer to lymphocytic host cells. There may also be mentioned the HSV-1 TK (herpesvirus type 1 thymidine kinase) gene promoter, the adenoviral MLP (major late promoter), in particular of human adenovirus type 2, the RSV (Rous Sarcoma Virus) LTR (long terminal repeat), the CMV (Cytomegalovirus) early promoter, and the PGK (phosphoglycerate kinase) gene promoter, for example, permitting expression in a large number of cell types.

As disclosed herein altering species tropism is demonstrated in BAV by replacement of the native fiber protein region with a heterologous mammalian fiber protein region. The present invention also encompasses replacement of one bovine serotype adenovirus fiber region with another bovine serotype adenovirus fiber region wherein said replacement is associated with altered bovine cell specificity.

Alternatively, targeting of a recombinant BAV vector to a particular cell type can be achieved by constructing recombinant hexon and/or fiber genes. The protein products of these genes are involved in host cell recognition; therefore, the genes can be modified to contain peptide sequences that will allow the virus to recognize alternative host cells.

Among genes of interest which are useable in the context of the present invention, there may be mentioned:

- genes coding for cytokines such as interferons and interleukins;
- genes encoding lymphokines;

- genes encoding tumor suppressors;

5 - genes involved in cardiovascular disease including, but not limited to, oncogenes; genes encoding growth factors including, but not limited to, fibroblast growth factor (FGF), vascular endothelial growth factor (VEGF), and nerve growth factor (NGF); *e-nos*, tumor suppressor genes including, but not limited to, the Rb (retinoblastoma) gene; lipoprotein lipase; superoxide dismutase (SOD); catalase; oxygen and free radical scavengers; apolipoproteins; and *pai-1* (plasminogen activator inhibitor-1);

10 - genes coding for cellular enzymes or those produced by pathogenic organisms; and

15 - suicide genes. The HSV-1 TK suicide gene may be mentioned as an example. This viral TK enzyme displays markedly greater affinity compared to the cellular TK enzyme for certain nucleoside analogues (such as acyclovir or gancyclovir). It converts them to monophosphorylated molecules, which can themselves be converted by cellular enzymes to nucleotide precursors, which are toxic. These nucleotide analogues can be incorporated into replicating DNA molecules, hence incorporation occurs chiefly in the DNA of dividing cells. This incorporation can result in specific destruction of dividing cells such as cancer cells.

20 This list is not restrictive, and other genes of interest may be used in the context of the present invention.

25 It is also possible that only fragments of nucleotide sequences of genes can be used (where these are sufficient to generate a protective immune response or a specific biological effect) rather than the complete sequence as found in the wild-type organism. Where available, synthetic genes or fragments thereof can also be used. However, the present invention can be used with a wide variety of genes, fragments and the like, and is not limited to those set out above.

In some cases the gene for a particular antigen can contain a large number of introns or can be from an RNA virus, in these cases a complementary DNA copy (cDNA) can be used.

5 In order for successful expression of the gene to occur, it can be inserted into an expression vector together with a suitable promoter including enhancer elements and polyadenylation sequences. A number of eucaryotic promoter and polyadenylation sequences which provide successful expression of foreign genes in mammalian cells and construction of expression cassettes, are known in the art, for example in U.S. Patent 5,151,267, the disclosures of which are incorporated herein by
10 reference. The promoter is selected to give optimal expression of immunogenic protein which in turn satisfactorily leads to humoral, cell mediated and mucosal immune responses according to known criteria.

The foreign protein produced by expression *in vivo* in a recombinant virus-infected cell may be itself immunogenic. More than one foreign gene can be inserted
15 into the viral genome to obtain successful production of more than one effective protein.

Thus with the recombinant viruses of the present invention, it is possible to provide protection against a wide variety of diseases affecting cattle, humans and other mammals. Any of the recombinant antigenic determinants or recombinant live
20 viruses of the invention can be formulated and used in substantially the same manner as described for antigenic determinant vaccines or live vaccine vectors.

The present invention also includes pharmaceutical compositions comprising a therapeutically effective amount of a recombinant adenovirus vector, recombinant adenovirus or recombinant protein, prepared according to the methods of the
25 invention, in combination with a pharmaceutically acceptable vehicle and/or an adjuvant. Such a pharmaceutical composition can be prepared and dosages determined according to techniques that are well-known in the art. The pharmaceutical compositions of the invention can be administered by any known

administration route including, but not limited to, systemically (for example, intravenously, intratracheally, intravascularly, intrapulmonarily, intraperitoneally, intranasally, parenterally, enterically, intramuscularly, subcutaneously, intratumorally or intracranially) or by aerosolization or intrapulmonary instillation. Administration can take place in a single dose or in doses repeated one or more times after certain time intervals. The appropriate administration route and dosage will vary in accordance with the situation (for example, the individual being treated, the disorder to be treated or the gene or polypeptide of interest), but can be determined by one of skill in the art.

The invention also encompasses a method of treatment, according to which a therapeutically effective amount of a BAV vector, recombinant BAV, or host cell of the invention is administered to a mammalian subject requiring treatment.

The antigens used in the present invention can be either native or recombinant antigenic polypeptides or fragments. They can be partial sequences, full-length sequences, or even fusions (e.g., having appropriate leader sequences for the recombinant host, or with an additional antigen sequence for another pathogen). The preferred antigenic polypeptide to be expressed by the virus systems of the present invention contain full-length (or near full-length) sequences encoding antigens. Alternatively, shorter sequences that are antigenic (i.e., encode one or more epitopes) can be used. The shorter sequence can encode a "neutralizing epitope," which is defined as an epitope capable of eliciting antibodies that neutralize virus infectivity in an *in vitro* assay. Preferably the peptide should encode a "protective epitope" that is capable of raising in the host a "protective immune response;" i.e., an antibody- and/or a cell-mediated immune response that protects an immunized host from infection.

The antigens used in the present invention, particularly when comprised of short oligopeptides, can be conjugated to a vaccine carrier. Vaccine carriers are well known in the art: for example, bovine serum albumin (BSA), human serum albumin

(HSA) and keyhole limpet hemocyanin (KLH). A preferred carrier protein, rotavirus VP6, is disclosed in EPO Pub. No. 0259149, the disclosure of which is incorporated by reference herein.

Genes for desired antigens or coding sequences thereof which can be inserted include those of organisms which cause disease in mammals, particularly bovine pathogens such as bovine rotavirus, bovine coronavirus, bovine herpes virus type 1, bovine respiratory syncytial virus, bovine parainfluenza virus type 3 (BPI-3), bovine diarrhea virus, *Pasteurella haemolytica*, *Haemophilus somnus* and the like. Genes encoding antigens of human pathogens also useful in the practice of the invention.

The vaccines of the invention carrying foreign genes or fragments can also be orally administered in a suitable oral carrier, such as in an enteric-coated dosage form. Oral formulations include such normally-employed excipients as, for example, pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharin cellulose, magnesium carbonate, and the like. Oral vaccine compositions may be taken in the form of solutions, suspensions, tablets, pills, capsules, sustained release formulations, or powders, containing from about 10% to about 95% of the active ingredient, preferably about 25% to about 70%. Oral and/or intranasal vaccination may be preferable to raise mucosal immunity (which plays an important role in protection against pathogens infecting the respiratory and gastrointestinal tracts) in combination with systemic immunity.

In addition, the vaccine can be formulated into a suppository. For suppositories, the vaccine composition will include traditional binders and carriers, such as polyalkaline glycols or triglycerides. Such suppositories may be formed from mixtures containing the active ingredient in the range of about 0.5% to about 10% (w/w), preferably about 1% to about 2%.

Protocols for administering to animals the vaccine composition(s) of the present invention are within the skill of the art in view of the present disclosure. Those skilled in the art will select a concentration of the vaccine composition in a

dose effective to elicit an antibody and/or T-cell mediated immune response to the antigenic fragment. Within wide limits, the dosage is not believed to be critical. Typically, the vaccine composition is administered in a manner which will deliver between about 1 to about 1,000 micrograms of the subunit antigen in a convenient volume of vehicle, e.g., about 1-10 cc. Preferably, the dosage in a single immunization will deliver from about 1 to about 500 micrograms of subunit antigen, more preferably about 5-10 to about 100-200 micrograms (e.g., 5-200 micrograms).

The timing of administration may also be important. For example, a primary inoculation preferably may be followed by subsequent booster inoculations if needed. It may also be preferred, although optional, to administer a second, booster immunization to the animal several weeks to several months after the initial immunization. To insure sustained high levels of protection against disease, it may be helpful to readminister a booster immunization to the animals at regular intervals, for example once every several years. Alternatively, an initial dose may be administered orally followed by later inoculations, or vice versa. Preferred vaccination protocols can be established through routine vaccination protocol experiments.

The dosage for all routes of administration of *in vivo* recombinant virus vaccine depends on various factors including, the size of patient, nature of infection against which protection is needed, carrier and the like and can readily be determined by those of skill in the art. By way of non-limiting example, a dosage of between 10^3 pfu and 10^{15} pfu, preferably between 10^5 and 10^{13} pfu, more preferably between 10^6 to 10^{11} pfu and the like can be used. As with *in vitro* subunit vaccines, additional dosages can be given as determined by the clinical factors involved.

In some embodiments of the invention, recombinant cell lines are produced by constructing an expression cassette comprising the BAV E1 region, and/or other essential gene region and transforming host cells therewith to provide complementing cell lines or cultures expressing the E1 proteins for use with replication-defective

bovine adenoviruses modified to have altered tropism and lacking E1 function. These recombinant complementing cell lines are capable of allowing a defective recombinant BAV with deleted E1 sequences to replicate and express a desired foreign gene or fragment thereof which is optionally encoded within the recombinant BAV. These cell lines are also extremely useful in generating recombinant BAV, having an E3 gene deletion replaced by heterologous nucleotide sequence encoding for a foreign gene or fragment, by *in vivo* recombination following DNA-mediated cotransfection. More generally, defective recombinant BAV vectors, lacking one or more essential functions encoded by the BAV genome, can be propagated in appropriate complementing cell lines, wherein a particular complementing cell line provides a function or functions that is (are) lacking in a particular defective recombinant BAV vector. Complementing cell lines can provide viral functions through, for example, co-infection with a helper virus, or by integrating or otherwise maintaining in stable form a fragment of a viral genome encoding a particular viral function.

In one embodiment of the invention, the recombinant expression cassette can be obtained by cleaving a BAV genome with an appropriate restriction enzyme to produce a DNA fragment representing the left end or the right end of the genome comprising E1 or E3 gene region sequences, respectively and inserting the left or right end fragment into a cloning vehicle, such as a plasmid, and thereafter inserting at least one heterologous DNA sequence into the E1 or E3 deletion with or without the control of an exogenous promoter. The recombinant expression cassette is contacted with a BAV genome within an appropriate cell and, through homologous recombination or other conventional genetic engineering method, a recombinant BAV genome is obtained. Appropriate cells include both prokaryotic cells, such as, for example, *E. coli*, and eukaryotic cells. Examples of suitable eukaryotic cells include, but are not limited to, MDBK cells, MDBK cells expressing adenovirus E1 function, primary fetal bovine retina cells, and cells expressing functions that are

equivalent to those of the previously-recited cells. Restriction fragments of the BAV genome other than those comprising the E1 or E3 regions are also useful in the practice of the invention and can be inserted into a cloning vehicle such that heterologous sequences may be inserted into non-E1 and E3 BAV sequences. These DNA constructs can then undergo recombination *in vitro* or *in vivo*, with a BAV genome, either before or after transformation or transfection of a suitable host cell as described above. For the purposes of the present invention, a BAV genome can be either a full-length genome or a genome containing a deletion in a region other than that deleted in the fragment with which it recombines, as long as the resulting recombinant BAV genome contains BAV sequences required for replication and packaging. Methods for transfection, cell culture and recombination in procaryotic and eukaryotic cells such as those described above are well-known to those of skill in the art.

In another embodiment of the invention, the function of any viral region which may be mutated or deleted in any particular viral vector can be supplied (to provide a complementing cell line) by co-infection of cells with a virus which expresses the function that the vector lacks.

If an insertion is made in a gene essential for viral replication, the adenovirus must be grown in an appropriate complementing cell line (*i.e.*, a helper cell line). In human adenoviruses, certain open reading frames in the E4 region (ORF 3 and ORF 6/7) are essential for viral replication. Deletions in analogous open reading frames in the E4 region of BAV-3 could necessitate the use of a helper cell line for growth of the viral vector.

The BAV E1 gene products of the adenovirus of the invention transactivate most of the cellular genes, and therefore, cell lines which constitutively express E1 proteins can express cellular polypeptides at a higher level than normal cell lines. The recombinant mammalian, particularly bovine, cell lines of the invention can be used to prepare and isolate polypeptides, including those such as (a) proteins

associated with adenovirus E1A proteins: e.g. p300, retinoblastoma (Rb) protein, cyclins, kinases and the like; (b) proteins associated with adenovirus E1B protein: e.g. p53 and the like; (c) growth factors, such as epidermal growth factor (EGF), transforming growth factor (TGF) and the like; (d) receptors such as epidermal growth factor receptor (EGF-R), fibroblast growth factor receptor (FGF-R), tumor necrosis factor receptor (TNF-R), insulin-like growth factor receptor (IGF-R), major histocompatibility complex class I receptor and the like; (e) proteins encoded by proto-oncogenes such as protein kinases (tyrosine-specific protein kinases and protein kinases specific for serine or threonine), p21 proteins (guanine nucleotide-binding proteins with GTPase activity) and the like; (f) other cellular proteins such as actins, collagens, fibronectins, integrins, phosphoproteins, proteoglycans, histones and the like, and (g) proteins involved in regulation of transcription such as TATA-box-binding protein (TBP), TBP-associated factors (TAFs), Sp1 binding protein and the like.

The invention also includes a method for providing gene delivery to a mammal, such as a bovine or a human or other mammal in need thereof, to control a gene deficiency, to provide a therapeutic gene or nucleotide sequence and/or to induce or correct a gene mutation. The method can be used, for example, in the treatment of conditions including, but not limited to hereditary disease, infectious disease, cardiovascular disease, and viral infection. The method comprises administering to said mammal a live recombinant bovine adenovirus comprising a modification in a capsid protein, or fragment thereof, wherein said capsid protein is associated with tropism and said modification is associated with altered tropism and wherein said adenovirus vector further comprises a foreign polynucleotide sequence encoding a non-defective form of said gene under conditions wherein the recombinant virus vector genome is incorporated into said mammalian genome or is maintained independently and extrachromosomally to provide expression of the required gene in the target organ or tissue. These kinds of techniques are currently

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being used by those of skill in the art for the treatment of a variety of disease conditions, non-limiting examples of which are provided above. Examples of foreign genes, nucleotide sequences or portions thereof that can be incorporated for use in a conventional gene therapy include, cystic fibrosis transmembrane conductance regulator gene, human minidystrophin gene, alpha-1-antitrypsin gene, genes involved in cardiovascular disease, and the like.

In particular, the practice of the present invention in regard to gene delivery in humans is intended for the prevention or treatment of diseases including, but not limited to, genetic diseases (for example, hemophilia, thalassemias, emphysema, Gaucher's disease, cystic fibrosis, Duchenne muscular dystrophy, Duchenne's or Becker's myopathy, *etc.*), cancers, viral diseases (for example, AIDS, herpesvirus infection, cytomegalovirus infection and papillomavirus infection), cardiovascular diseases, and the like. For the purposes of the present invention, the vectors, cells and viral particles prepared by the methods of the invention may be introduced into a subject either *ex vivo*, (*i.e.*, in a cell or cells removed from the patient) or directly *in vivo* into the body to be treated.

The following examples are provided to illustrate but not limit the invention.

EXAMPLES

Example 1: Construction of BAV600 containing a human fiber gene

To generate an BAV-3 vector with an altered tropism, the chimeric fiber gene construct containing the HAV-5 fiber knob fused to the BAV-3 tail and shaft was incorporated into the BAV-3 genome of BAV304, described in Reddy *et al.*, *supra* 1999 (Fig. 3). For the precise replacement of the wild-type BAV-3 fiber gene, a previously made plasmid pBAV301.gfp (Reddy *et al.*, 1999) was used for modification of BAV-3 fiber. The resulting transfer vector pBAV-301.G5FK contained a CMV promoter driven green fluorescent protein (GFP) expression

cassette inserted into the E3 region, the chimeric BAV-3/HAV-5 fiber gene, and E4. This transfer vector was used for incorporation of GFP cassette and modified fiber gene into the backbone of an E3 deleted BAV-3 infectious plasmid, p.FBAV302 (Zakhartchouk *et al.*, 1998), via homologous recombination in *E. coli* BJ5183 (Chartier *et al.*, 1996), creating plasmid pFBAV-600. The viral genome was released from the plasmid by PacI digestion and used to transfect cell line ATCC accession number PTA156; fetal bovine retinal cells expressing E1 protein (see Reddy *et al.* 1999, *supra*). The corresponding chimeric virus BAV600 was produced 21 days following transfection.

Example 2: Characterization of BAV600

BAV600 obtained from the transfection of fetal bovine retinal cells expressing E1 protein, ATCC accession number PTA156, was amplified in MDBK cells, and the viral DNA was extracted from infected cells. The DNA was analyzed after digestion with restriction enzyme *Bgl*/II and agarose gel electrophoresis (Figure 5A). As shown in Figures 5A-5B, both the parental BAV302 and BAV304 had *Bgl*/II fragment of 5.4 kb at the right end of viral genome. The HAV-5 fiber knob region introduces an additional *Bgl*/II restriction enzyme site within the BAV600 genome. Therefore, diagnostic 1.5 and 3.9 kb fragments were found after *Bgl*/II digestion. Southern blot analysis with the HAV-5 fiber knob probe demonstrated the expected hybridization pattern for *Bgl*/II-digested BAV600 (Figure 5B).

Expression and assembly of the chimeric BAV-3 and HAV-5 fiber protein by recombinant BAV600 were examined by immunoprecipitation assay. Metabolically radiolabeled immunoprecipitates from the parental (BAV304; Reddy *et al.*, 1999, *supra*) and chimeric (BAV600) viruses-infected MDBK cell lysates were subjected to SDS-PAGE under denaturing conditions. A wild-type HAV-5 containing a full-length fiber was also analyzed. Immunoprecipitation assay was carried out with a rabbit polyclonal antibody specific for the BAV3 fiber knob and an antifiber

monoclonal antibody, ID6.14. The ID6.14 antibody recognizes a trimerized HAV-5 fiber knob and neutralizes HAV-5 through binding to knob domain (Douglas *et al.*, 1996). As shown in Figure 6, the BAV-3 and BAV304 viruses contain fiber proteins with sizes of approximately 100 kDa which react with the rabbit polyclonal antibody specific for the BAV3 fiber knob, while the HAV-5 contains a fiber protein with a size of approximately 64 kDa. The presence of the HAV-5 fiber knob within the BAV600 chimeric virus was confirmed by immunoprecipitation analysis with the monoclonal antibody ID6.14 specific for the HAV-5 knob.

The biological titer of the fiber chimeric virus BAV600 was compared with the BAV-3 and parental virus BAV304. Biological titers determined with MDBK cell monolayers indicated maximum plaque-forming titers of 10^8 , 10^6 , and 10^5 PFU/ml for the BAV-3, BAV304, and BAV600, respectively. The result suggested that the fiber modification and GFP insertion in E3 region significantly alter the cellular production of the virus.

Example 3: Transduction of human cell lines by BAV600

To characterize the transduction efficiency of BAV304 and BAV600 in different human cell lines, FACS analysis was performed to determine the percentage of transduction of each cell line at different virus input (Fig 7A). Cells grown in T25 flasks were infected at an MOI of 1 and 5 with either BAV304 or BAV600. Forty-eight hours after infection, the percentage of GFP-fluorescence positive cells were determined by flow cytometry. The percentage of transduction of each cell line was quantitated, and the fraction of dose is shown in Figure 7B. 293 cells were equally susceptible to transduction with both viruses (indicating that both the HAV-5 and BAV-3 receptors are present on the cell surface.) The transduction of HeLa and HEp-2 cells with BAV304 is dose dependent, with about 6% and 1% respectively at an MOI of 1 and about 25% and 5% respectively at an MOI of 5. Both cells were efficiently transduced with BAV600. The percentage of transduction with BAV600

reaches maximum level even at an MOI of 1 (94% and 93% for HeLa and Hep-2 respectively). In contrast less-efficient transduction of A549 cells with BAV600 was observed. These data taken together demonstrate that the BAV600 containing HAV-5 fiber knob was clearly superior to the BAV304 vector in transduction of human cell lines.

Example 4: HAV-5 and BAV-3 neutralizing antibodies in human serum

Preexisting neutralizing antibodies against HAV-5 in clinical patients represent a major obstacle for efficient use of HAV-5 in human gene therapy protocol. In order to explore the possibility for use of BAV-3 as an alternative vector to HAV-5-derived vectors, it was determined whether preexisting anti-HAV-5 neutralizing antibodies were also cross-reactive with BAV-3. 105 random samples of human sera from clinical patients were tested. Three (#50, 97, and 102) were found containing high titer of HAV-5 neutralizing antibodies ranging between 1:800 to 1,6000. These sera were tested for their ability to inhibit BAV-3-induced plaque formation on MDBK cells. Our data demonstrated that none of these HAV-5 positive sera showed effect on BAV-3-induced plaque formation at a dilution of 1/50.

Example 5: Replication of BAV-3 in human cell lines

Virus production and the time course of virus infection were studied in different human cell lines to determine their degree of permissivity for BAV-3 growth. Confluent monolayer cultures of each cell line (HeLa, HEp-2, 293 and A549) were infected-with BAV-3 at an MOI of 10 and virus production at different time intervals after infection was assayed by titration of the cell lysates on MDBK cell monolayers. Virus growth in permissive MDBK cells resulted in, as expected, maximum yields of 10^8 pfu/ml by 48 hours after infection. In contrast, the level of

BAV-3 production in all four human cell lines was constantly diminished, suggesting that there is a complete absence of viral replication in these human cell lines.

Example 6: Expression of early and later BAV-3 proteins in human cell lines

5 Viral proteins include early proteins (E1B small and single-stranded DNA binding protein [DBP]) and late proteins (penton base and fiber). To identify the expression of early and late viral proteins in human cell lines, viral protein production was analyzed by Western immunoblotting. Cultures were infected with BAV-3 at an MOI of 10. At intervals after infection, cell extracts were prepared from each culture, 10 separated on 10% SDS-PAGE, and transferred to nitrocellulose. Antigens immobilized on the nitrocellulose sheets were probed by reaction with rabbit polyclonal antibodies against E1B small, DBP, penton base, and fiber respectively. As expected, the E1B small and DBP antisera reacted with bands in 19 and 50 kDa, respectively, from BAV-3-infected MDBK cells. In contrast, all human cell lines 15 except 293 cell lines showed no positive reactions with anti-E1B small or DBP polyclonal antibodies. No structural proteins were detected from BAV-3- infected human cell lines. These results indicated that the replication of BAV-3 in the majority of human cells tested in this study was blocked at E1B small level.

20 *Example 7: Neutralization of BAV600 by an monoclonal antibody specific for HAV-5 fiber knob*

 It was hypothesized that BAV600 carrying the HAV-5 fiber knob should be neutralized by an antibody specific for HAV-5 knob. To confirm this, duplicate aliquots containing 100 pfu of BAV-3 or BAV600 were incubated at room 25 temperature for two hours with serial twofold dilutions of a rabbit polyclonal antibody specific for the BAV3 fiber knob or a monoclonal antibody, 1D6.14, against HAV-5 fiber knob domain. MDBK cells were then infected with pre-incubated BAV-3 or BAV600 virus. Cells were incubated for 14 days to allow a complete CPE

to develop. The data show that that none of the viruses were neutralized by serum from normal rabbit serum or a control monoclonal antibody 2C8 specific for bovine herpesvirus gD protein. BAV-3 and BAV600 were each neutralized by a rabbit polyclonal antibody specific for the BAV3 fiber knob (1:800) and ID6.14 (1:3,200), respectively. However, neither virus was neutralized by the reciprocal antiserum even at a dilution of 1:50. This further confirmed that BAV600 carried the HAV-5 fiber knob.

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